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STRUCTURE FILE UPDATES: 22 FEB 2006 HIGHEST RN 874945-83-2
DICTIONARY FILE UPDATES: 22 FEB 2006 HIGHEST RN 874945-83-2

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*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added, *
* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

Structure search iteration limits have been increased. See HELP SLIMITS
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=> file caplus
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FILE COVERS 1907 - 24 Feb 2006 VOL 144 ISS 10
FILE LAST UPDATED: 23 Feb 2006 (20060223/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply.

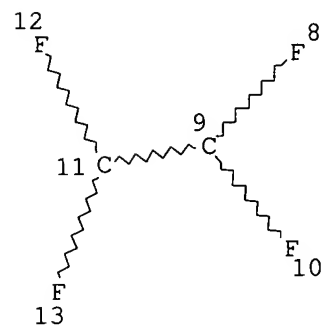
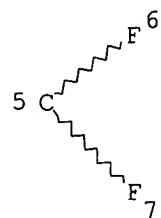
They are available for your review at:

<http://www.cas.org/infopolicy.html>

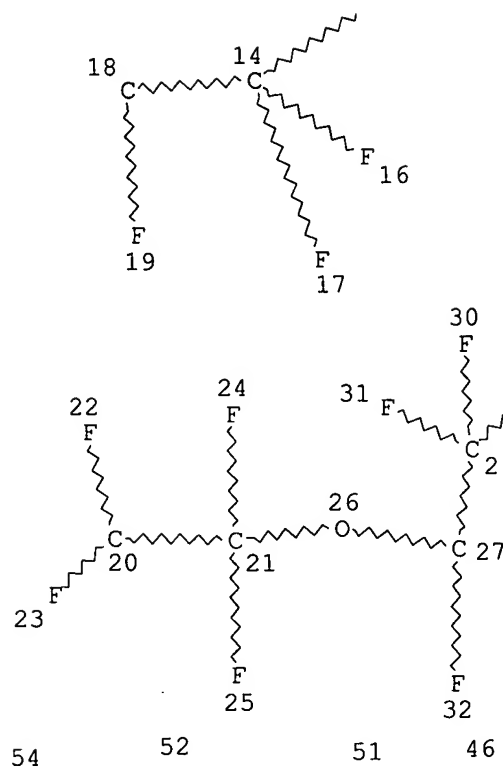
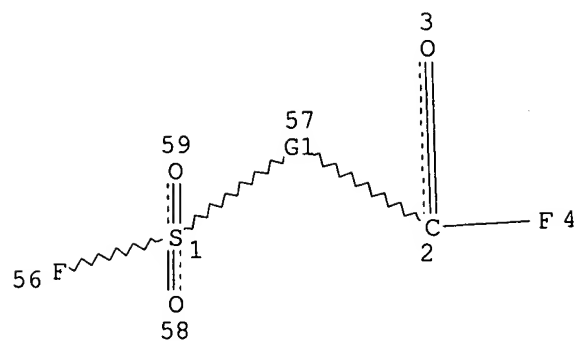
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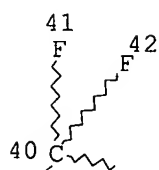
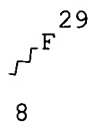
L18 STR



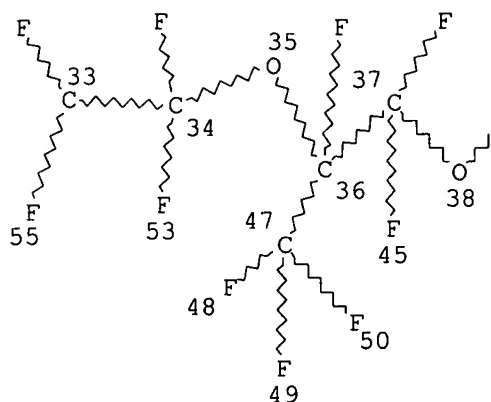
Page 1-A



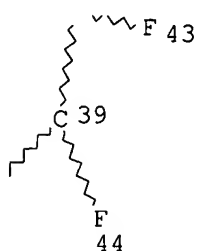
Page 2-A



Page 2-B



Page 3-A



Page 3-B

VAR G1=5-1 5-2/11-1 9-2/18-1 18-2/20-1 27-2/33-1 39-2

NODE ATTRIBUTES:

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 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 58 59

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 59

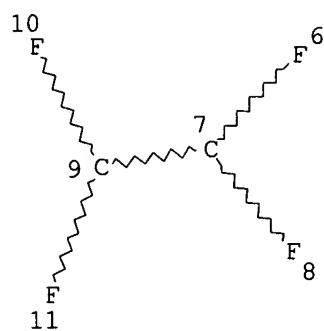
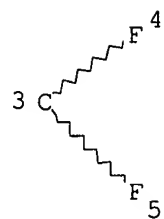
STEREO ATTRIBUTES: NONE

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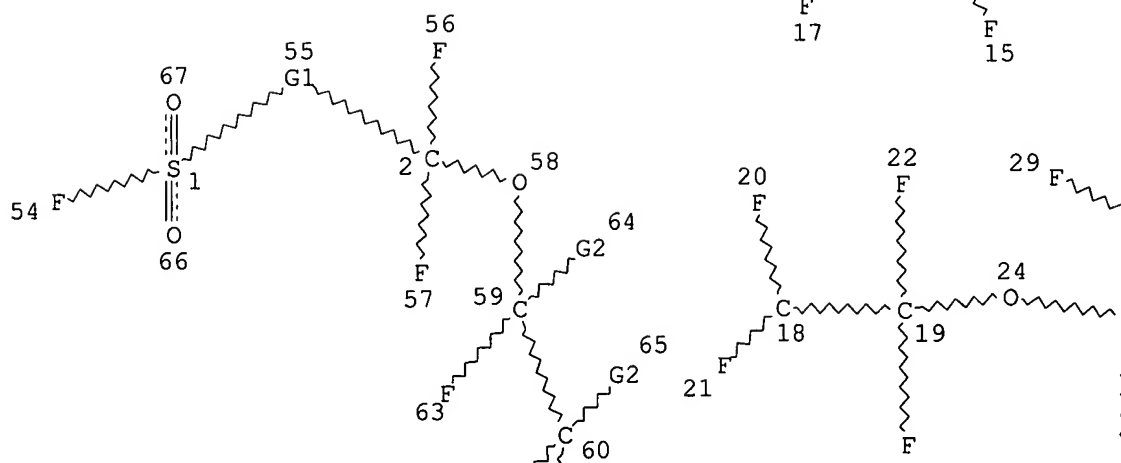
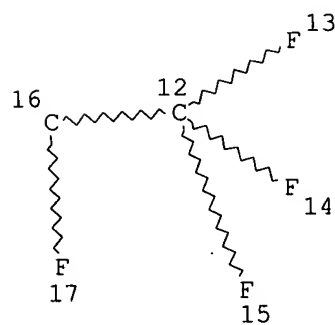
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 RGT)/RL

L23 STR

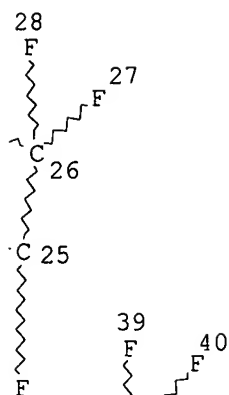
Cl 68Br 69



Page 1-A



Page 2-A



The chemical structure is a complex, branched molecule with multiple fluorinated side chains. The atoms are numbered as follows: 23, 3, 49, 35, 33, 50, 52, 62, 61, 31, 32, 51, 53, 45, 46, 47, 48, 43, 34, and 3. The structure features a central carbon-carbon bond (C31-C32) and several ether linkages (O33, O34). The fluorinated chains are represented by zigzag lines, and the terminal fluorine atoms are labeled 'F'.

A chemical structure diagram of a branched polymer. The main chain consists of two carbon atoms, labeled 38 and 37, connected by a single bond. Carbon 38 is bonded to a fluorine atom (F0) and a chain segment labeled 44. Carbon 37 is bonded to a fluorine atom (F42) and a chain segment labeled 36. The chain segment 44 is further branched, with a fluorine atom (F41) and another chain segment labeled 41. The chain segment 36 is also branched, with a fluorine atom (F42) and another chain segment labeled 42.

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DEFAULT MLEVEL IS ATOM

MLEVEL IS CLASS AT 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17
 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38
 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 56 57 58 59 60
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DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 69

STEREO ATTRIBUTES: NONE

L25 6 SEA FILE=REGISTRY SSS FUL L23

L27 18 SEA FILE=CAPLUS ABB=ON PLU=ON L25 (L) PREP/RL

L28 8 SEA FILE=CAPLUS ABB=ON PLU=ON L22 AND L27

=> d ibib abs hitind hitstr L28 1-8

L28 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:753175 CAPLUS

DOCUMENT NUMBER: 141:260266

TITLE: Process for preparing (per)fluorohalogen ethers by the
 reaction of acyl fluorides with halogenated
 1,2-difluoroethylenes

INVENTOR(S): Tortelli, Vito; Calini, Pierangelo; Millefanti,
 Stefano

PATENT ASSIGNEE(S): Solvay Solexis S.p.A., Italy

SOURCE: Eur. Pat. Appl., 8 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1457484	A1	20040915	EP 2004-4344	20040226
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2004269535	A2	20040930	JP 2004-65994	20040309
US 2004199009	A1	20041007	US 2004-795995	20040310
CN 1539818	A	20041027	CN 2004-10033085	20040311
PRIORITY APPLN. INFO.:			IT 2003-MI444	A 20030311

OTHER SOURCE(S): CASREACT 141:260266; MARPAT 141:260266

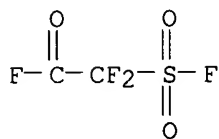
AB A process for preparing (per)fluorohalogen ethers containing the sulfonyl
 fluoride group FSO₂RCF₂OCAFCA1F₂ [A, A1 = Cl, Br; R = (per)fluorinated
 optionally containing one or more oxygen atoms] is described which comprises
 the reaction of acyl fluorides FSO₂RCOF in the liquid phase with elemental
 fluorine and with olefinic compds. CAF:CA1F at -120° to
 -20°, optionally in the presence of a solvent inert under the
 reaction conditions.

IC ICM C07C303-22

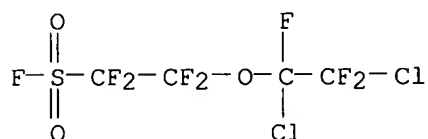
ICS C07C309-82

CC 23-12 (Aliphatic Compounds)

Section cross-reference(s): 45
 IT 76-15-3, cfc 115 359-21-7 598-88-9, 1,2-Dichloro-1,2-difluoroethylene
 677-67-8
 RL: **RCT (Reactant); RACT (Reactant or reagent)**
 (process for preparing (per)fluorohalogen ethers by the reaction of acyl
 fluorides with halogenated 1,2-difluoroethylenes)
 IT 144728-59-6P
 RL: **SPN (Synthetic preparation); PREP (Preparation)**
 (process for preparing (per)fluorohalogen ethers by the reaction of acyl
 fluorides with halogenated 1,2-difluoroethylenes)
 IT 677-67-8
 RL: **RCT (Reactant); RACT (Reactant or reagent)**
 (process for preparing (per)fluorohalogen ethers by the reaction of acyl
 fluorides with halogenated 1,2-difluoroethylenes)
 RN 677-67-8 CAPLUS
 CN Acetyl fluoride, difluoro(fluorosulfonyl)- (6CI, 7CI, 8CI, 9CI) (CA INDEX
 NAME)



IT 144728-59-6P
 RL: **SPN (Synthetic preparation); PREP (Preparation)**
 (process for preparing (per)fluorohalogen ethers by the reaction of acyl
 fluorides with halogenated 1,2-difluoroethylenes)
 RN 144728-59-6 CAPLUS
 CN Ethanesulfonyl fluoride, 2-(1,2-dichloro-1,2,2-trifluoroethoxy)-1,1,2,2-
 tetrafluoro- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L28 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN
 ACCESSION NUMBER: 2004:668857 CAPLUS
 DOCUMENT NUMBER: 142:59591
 TITLE: Synthesis of 3,6-dioxo-Δ⁷-4-trifluoromethyl
 perfluorooctyl trifluoromethyl sulfonimide:
 bis[(perfluoroalkyl)sulfonyl] superacid monomer and
 polymer
 AUTHOR(S): Thomas, Brian H.; Shafer, Gregory; Ma, Jing Ji; Tu,
 Ming-Hu; DesMarteau, Darryl D.
 CORPORATE SOURCE: H.L. Hunter Hall Chemistry Laboratory, Chemistry
 Department, Clemson University, Clemson, SC,
 29634-1905, USA
 SOURCE: Journal of Fluorine Chemistry (2004), 125(8),
 1231-1240

CODEN: JFLCAR; ISSN: 0022-1139
 PUBLISHER: Elsevier B.V.
 DOCUMENT TYPE: Journal
 LANGUAGE: English

AB A new type of ion exchange polymer, bis[(perfluoroalkyl)sulfonyl]imide ionomers (PFSI), were developed by the copolymn. of sodium 3,6-dioxa-Δ7-4-trifluoromethyl perfluorooctyl trifluoromethyl sulfonimide with tetrafluoroethylene (TFE) using an aqueous redox initiation system in an emulsion type polymerization. These polymers were prepared in various

equivalent wts. and processed into functional membranes. The new ionomers exhibit excellent chemical and thermal stability. The materials have high potential for electrochem. applications especially as solid polymer electrolytes

(SPE) in proton exchange membrane (PEM) fuel cells.

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
 Section cross-reference(s): 35, 38

IT **677-67-8P**, Fluorosulfonyldifluoroacetyl fluoride
 RL: PEP (Physical, engineering or chemical process); PRP (Properties); PUR (Purification or recovery); PYP (Physical process); **RCT (Reactant)**; SPN (Synthetic preparation); PREP (Preparation); PROC (Process); **RACT (Reactant or reagent)**

(compound 4; synthesis of 3,6-dioxa-Δ7-4-trifluoromethyl perfluorooctyl trifluoromethyl sulfonimide, bis[(perfluoroalkyl)sulfonyl] superacid monomer and polymer)

IT 64346-22-1P **78010-39-6P**
 RL: PEP (Physical, engineering or chemical process); PRP (Properties); PUR (Purification or recovery); PYP (Physical process); RCT (Reactant); SPN (Synthetic preparation); **PREP (Preparation)**; PROC (Process); **RACT (Reactant or reagent)**

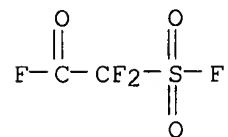
(compound 9; synthesis of 3,6-dioxa-Δ7-4-trifluoromethyl perfluorooctyl trifluoromethyl sulfonimide, bis[(perfluoroalkyl)sulfonyl] superacid monomer and polymer)

IT **677-67-8P**, Fluorosulfonyldifluoroacetyl fluoride
 RL: PEP (Physical, engineering or chemical process); PRP (Properties); PUR (Purification or recovery); PYP (Physical process); **RCT (Reactant)**; SPN (Synthetic preparation); PREP (Preparation); PROC (Process); **RACT (Reactant or reagent)**

(compound 4; synthesis of 3,6-dioxa-Δ7-4-trifluoromethyl perfluorooctyl trifluoromethyl sulfonimide, bis[(perfluoroalkyl)sulfonyl] superacid monomer and polymer)

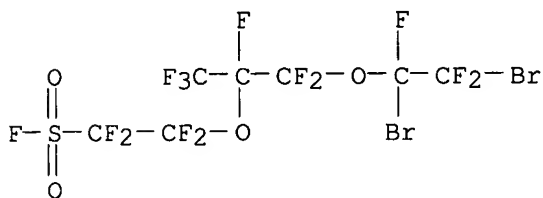
RN 677-67-8 CAPLUS

CN Acetyl fluoride, difluoro(fluorosulfonyl)- (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)



IT **78010-39-6P**
 RL: PEP (Physical, engineering or chemical process); PRP (Properties); PUR (Purification or recovery); PYP (Physical process); RCT (Reactant); SPN (Synthetic preparation); **PREP (Preparation)**; PROC (Process); **RACT (Reactant or reagent)**
 (compound 9; synthesis of 3,6-dioxa-Δ7-4-trifluoromethyl

perfluorooctyl trifluoromethyl sulfonimide,
bis[(perfluoroalkyl)sulfonyl] superacid monomer and polymer)
RN 78010-39-6 CAPLUS
CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 58 THERE ARE 58 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L28 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:289553 CAPLUS

DOCUMENT NUMBER: 140:321901

TITLE: Unsaturated fluorohydrocarbyl fluoroalkylsulfonates as substitutes for unsaturated fluoroalkylsulfonyl fluorides, and their manufacture

INVENTOR(S): Uematsu, Nobuyuki; Hoshi, Nobuto; Koga, Takehiro; Gronvald, Oliver; Ikeda, Masanori

PATENT ASSIGNEE(S): Asahi Kasei Corporation, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 18 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2004107313	A2	20040408	JP 2002-350246	20021202
PRIORITY APPLN. INFO.:			JP 2002-215050	A 20020724
OTHER SOURCE(S):		MARPAT 140:321901		

AB The fluorosulfonates, useful as monomers for separators for fuel cells and electrolysis of NaCl, etc., are CF₂:CF[OCF₂CF(CF₃)]_nO(CF₂)_mSO₃Rf (I; Rf = fluorohydrocarbyl, m = 1-5; n = 0-2). Thus, CF₂:CFOCF₂CF₂SO₃H was treated with CH₂:CF₂ to give I (Rf = CF₂Me, m = 2, n = 0).

IC ICM C07C309-10

ICS C07C303-28; C08F016-30

CC 35-2 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 23, 52, 72

IT **78010-39-6P** 111173-24-1P 677315-21-8P 677315-22-9P
677315-24-1P 677315-25-2P 677315-27-4P 677315-28-5P 677315-31-0P
677315-32-1P 677315-33-2P 677315-34-3P

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP**

(Preparation); RACT (Reactant or reagent)

(manufacture of unsatd. fluorohydrocarbyl fluoroalkylsulfonates as monomers for separators for fuel cells and electrolysis of NaCl)

IT 75-38-7, Vinylidene fluoride 75-89-8, 2,2,2-Trifluoroethanol 76-37-9
920-66-1 **4089-57-0** 16090-14-5 26953-98-0

RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of unsatd. fluorohydrocarbyl fluoroalkylsulfonates as monomers for separators for fuel cells and electrolysis of NaCl)

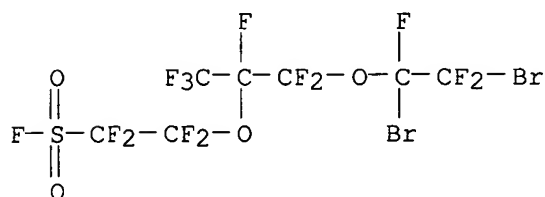
IT 78010-39-6P

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP** (Preparation); RACT (Reactant or reagent)

(manufacture of unsatd. fluorohydrocarbyl fluoroalkylsulfonates as monomers for separators for fuel cells and electrolysis of NaCl)

RN 78010-39-6 CAPLUS

CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



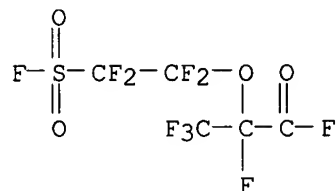
IT 4089-57-0

RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of unsatd. fluorohydrocarbyl fluoroalkylsulfonates as monomers for separators for fuel cells and electrolysis of NaCl)

RN 4089-57-0 CAPLUS

CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]- (9CI) (CA INDEX NAME)



L28 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2003:163674 CAPLUS

DOCUMENT NUMBER: 138:169855

TITLE: Process for the synthesis of perfluorosulfonylalkyl hypofluorites

INVENTOR(S): Navarrini, Walter

PATENT ASSIGNEE(S): Ausimont S.p.A., Italy

SOURCE: Ital. Appl., 25 pp.

CODEN: ITXXCZ

DOCUMENT TYPE: Patent

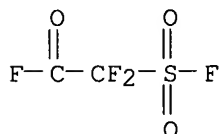
LANGUAGE: Italian

FAMILY ACC. NUM. COUNT: 1

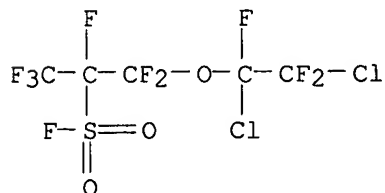
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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IT 2000MI1846	A1	20020208	IT 2000-MI1846	20000808
IT 1318672	B1	20030827		

PRIORITY APPLN. INFO.: IT 2000-MI1846 20000808
 OTHER SOURCE(S): CASREACT 138:169855; MARPAT 138:169855
 AB Hypofluorites FSO₂-Rf-CF₂OF [Rf = CF₂, CF₂CF₂, CF(CF₃), CF₂CF₂OCF(CF₃)] were prepared by fluorination of acyl fluorides FSO₂-Rf-COF or corresponding sultones [when Rf = CF₂, OCF(CF₃)] over a supported CsF or KF catalyst. Thus, fluorination of perfluoropropene sultone (2 mmol) with 4 mmol F₂ over a CsF/NaF catalyst (1 h at 200 mbar and room temperature) yielded FSO₂CF(CF₃)CF₂OF which reacted with 8 mmol CFCl:CFCl to afford 53% FSO₂CF(CF₃)CF₂OCFClCF₂Cl.
 IC ICM C07C309-78
 CC 23-11 (Aliphatic Compounds)
 IT 74-85-1, Ethylene, reactions 75-01-4, Chloroethylene, reactions
 79-38-9, 2 Chloro 1 1 2 trifluoroethylene 540-59-0, 1 2 Dichloroethylene
 598-88-9, 1 2 Dichloro 1 2 difluoroethylene 677-67-8 697-18-7
 773-15-9 89413-95-6 89413-97-8
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of perfluorosulfonylalkyl hypofluorites from perfluorosulfonylalkanoyl fluorides)
 IT 115784-53-7P 144728-64-3P 496922-45-3P 496922-46-4P
 496922-47-5P 496922-48-6P 496922-49-7P 496922-50-0P 496922-51-1P
 496922-52-2P 496922-54-4P 496922-55-5P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of perfluorosulfonylalkyl hypofluorites from perfluorosulfonylalkanoyl fluorides)
 IT 677-67-8
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of perfluorosulfonylalkyl hypofluorites from perfluorosulfonylalkanoyl fluorides)
 RN 677-67-8 CAPLUS
 CN Acetyl fluoride, difluoro(fluorosulfonyl)- (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)



IT 144728-64-3P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of perfluorosulfonylalkyl hypofluorites from perfluorosulfonylalkanoyl fluorides)
 RN 144728-64-3 CAPLUS
 CN 2-Propanesulfonyl fluoride, 1-(1,2-dichloro-1,2,2-trifluoroethoxy)-1,1,2,3,3,3-hexafluoro- (9CI) (CA INDEX NAME)



L28 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:615562 CAPLUS
 DOCUMENT NUMBER: 137:169968
 TITLE: Manufacture of perfluorovinyl ether monomer having
 sulfonamide group and its use for solid electrolyte
 membrane
 INVENTOR(S): Ikeda, Masanori; Hoshi, Nobuto; Uematsu, Nobuyuki;
 Koga, Takehiro
 PATENT ASSIGNEE(S): Asahi Kasei Kabushiki Kaisha, Japan
 SOURCE: PCT Int. Appl., 215 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002062749	A1	20020815	WO 2002-JP854	20020201
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1359142	A1	20031105	EP 2002-711282	20020201
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
CN 1500075	A	20040526	CN 2002-807780	20020201
US 2004122256	A1	20040624	US 2003-470802	20030801
PRIORITY APPLN. INFO.:			JP 2001-25018	A 20010201
			JP 2001-30955	A 20010207
			JP 2001-278418	A 20010913
			JP 2001-342172	A 20011107
			JP 2001-343780	A 20011108
			JP 2001-343931	A 20011108
			WO 2002-JP854	W 20020201

OTHER SOURCE(S): MARPAT 137:169968

AB A perfluorovinyl ether monomer represented by
 $\text{CF}_2\text{CF}(\text{OCF}_2\text{CFCF}_3)\text{mO}(\text{CF}_2)_n\text{SO}_2\text{NR}_1\text{R}_2$ (wherein m = 0-5 integer; n = 1-5 integer; R₁, R₂ = H, C₁-10 (un)substituted hydrocarbonyl, substituted silyl; R₁ and R₂ may be bonded to each other to form a ring) and its polymers are prepared and the polymer films are used as solid electrolyte membrane. Neutralization of $\text{CF}_3\text{CF}(\text{COF})\text{OCF}_2\text{CF}_2\text{SO}_3\text{F}$ with Na_2CO_3 , amidation with diethylamine and n-BuLi, and decarboxylation gave $\text{CF}_2\text{:CFOCF}_2\text{CF}_2\text{SO}_3\text{Net}_2$. Copolymn. of this monomer with tetrafluoroethylene and press molding at 250° gave a membrane useful for solid electrolyte.

IC ICM C07C311-24

ICS C07C303-36; C07F007-12; C08F214-26; C08F216-14; H01M008-02

CC 35-2 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 38, 52

IT 75549-02-9P 75718-06-8P 78010-39-6P 144728-59-6P

445293-56-1P 445293-57-2P 445293-58-3P 445293-59-4P 445293-60-7P

445293-61-8P 446312-49-8P 446312-51-2P 446312-52-3P 446312-53-4P

446312-54-5P 446312-55-6P 446312-56-7P 446312-57-8P 446312-58-9P

446312-59-OP 446312-61-4P 446312-62-5P 446312-63-6P 446312-65-8P
 446312-68-1P 446312-69-2P 446312-70-5P 446312-71-6P 446312-72-7P
 446312-75-OP

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP**
(Preparation); RACT (Reactant or reagent)

(manufacture of perfluorovinyl ether monomer having sulfonamide group for preparation of solid electrolyte membrane)

IT 62-53-3, Aniline, reactions 75-64-9, tert-Butylamine, reactions
 109-89-7, Diethylamine, reactions 109-97-7, Pyrrole 124-40-3,
 Dimethylamine, reactions 288-32-4, Imidazole, reactions 999-97-3,
 Hexamethyldisilazane 1070-89-9, Sodium hexamethyldisilazide **4089-57**
-0 4089-58-1 29514-94-1 77545-08-5

RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of perfluorovinyl ether monomer having sulfonamide group for preparation of solid electrolyte membrane)

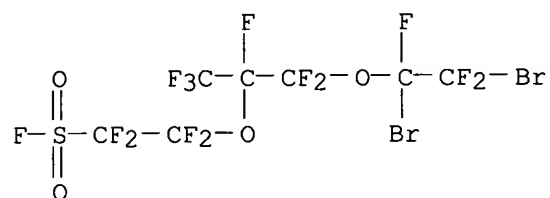
IT **78010-39-6P 144728-59-6P**

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP**
(Preparation); RACT (Reactant or reagent)

(manufacture of perfluorovinyl ether monomer having sulfonamide group for preparation of solid electrolyte membrane)

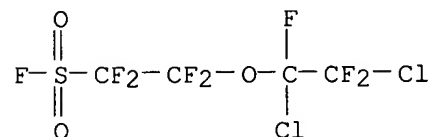
RN 78010-39-6 CAPLUS

CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



RN 144728-59-6 CAPLUS

CN Ethanesulfonyl fluoride, 2-(1,2-dichloro-1,2,2-trifluoroethoxy)-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



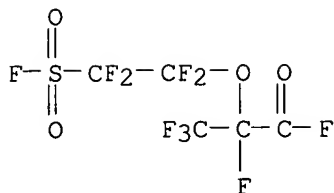
IT **4089-57-0 4089-58-1**

RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of perfluorovinyl ether monomer having sulfonamide group for preparation of solid electrolyte membrane)

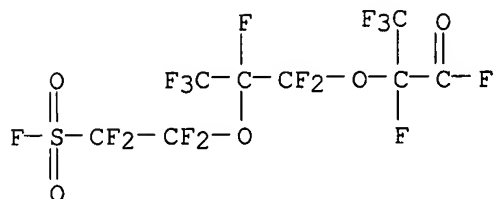
RN 4089-57-0 CAPLUS

CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]- (9CI) (CA INDEX NAME)



RN 4089-58-1 CAPLUS

CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,3,3,3-hexafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]propoxy]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L28 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:607663 CAPLUS

DOCUMENT NUMBER: 137:155315

TITLE: One-step manufacture of sulfonic acid group-containing fluoropolymers

INVENTOR(S): Koga, Takehiro; Hoshi, Nobuto; Ikeda, Masanori

PATENT ASSIGNEE(S): Asahi Kasei Corporation, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2002226514	A2	20020814	JP 2001-30967	20010207
PRIORITY APPLN. INFO.:			JP 2001-30967	20010207

AB The fluoropolymers, useful for fuel cell electrolytes, are manufactured by acid treatment of polymers having repeating units $\text{CF}_2\text{CF}[(\text{OCF}_2\text{CF}(\text{CF}_3))_m\text{O}(\text{CF}_2)_n\text{SO}_2\text{NR}_1\text{R}_2]$ ($\text{R}_1, 2 = \text{H}, \text{alkyl}, \text{aryl}, \text{aralkyl}, \text{silyl}$; $\text{R}_1\text{-R}_2$ may form ring; $m = 0, 1$; $n = 2, 3$). Sulfonamide groups-containing fluoropolymers, having repeating units $\text{CF}_2\text{CF}[\text{O}(\text{CF}_2)_n\text{SO}_2\text{NRR}_2]$ ($\text{R}_1, 2, m, n = \text{same as above}$), are also claimed. Thus, a fluoropolymer film having a unit $\text{CF}_2\text{CF}[(\text{OCF}_2\text{CF}(\text{CF}_3))_m\text{O}(\text{CF}_2)_n\text{SO}_2\text{NRR}_2]$, showing good antiblocking properties, was immersed in 3N H_2SO_4 at 130° for 1.5 h to convert SO_2NRR_2 to SO_3H .

IC ICM C08F008-12

ICS C08F016-30; H01M008-02

CC 35-8 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 52

IT 78010-39-6P 445293-56-1P 445293-59-4P 445293-60-7P

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP**

(Preparation); RACT (Reactant or reagent)

(one-step manufacture of sulfonic acid group-containing fluoropolymers by

acid

hydrolysis of sulfonamide group-containing precursors)

IT **4089-57-0**

RL: RCT (Reactant); RACT (Reactant or reagent)

(one-step manufacture of sulfonic acid group-containing fluoropolymers by

acid

hydrolysis of sulfonamide group-containing precursors)

IT **78010-39-6P**

RL: IMF (Industrial manufacture); RCT (Reactant); **PREP**

(Preparation); RACT (Reactant or reagent)

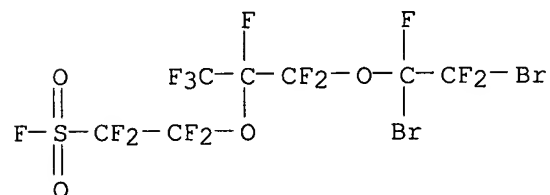
(one-step manufacture of sulfonic acid group-containing fluoropolymers by

acid

hydrolysis of sulfonamide group-containing precursors)

RN 78010-39-6 CAPLUS

CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



IT **4089-57-0**

RL: RCT (Reactant); RACT (Reactant or reagent)

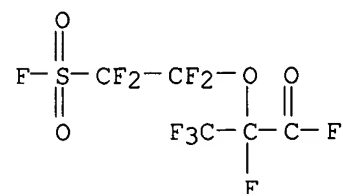
(one-step manufacture of sulfonic acid group-containing fluoropolymers by

acid

hydrolysis of sulfonamide group-containing precursors)

RN 4089-57-0 CAPLUS

CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]- (9CI) (CA INDEX NAME)



L28 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1982:617423 CAPLUS

DOCUMENT NUMBER: 97:217423

TITLE: Solutions of sulfonyl fluorides and fluoropolymers

INVENTOR(S): Silva, Raimund H.; Resnick, Paul R.; Smith, Roger A.

PATENT ASSIGNEE(S): du Pont de Nemours, E. I., and Co. , USA

SOURCE: U.S., 10 pp. Cont.-in-part of U.S. Ser. No. 79,173, abandoned.

DOCUMENT TYPE: CODEN: USXXAM
 LANGUAGE: Patent
 FAMILY ACC. NUM. COUNT: English
 PATENT INFORMATION: 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 4348310	A	19820907	US 1980-176595	19800808
JP 56050947	A2	19810508	JP 1980-131781	19800924
FR 2465753	A1	19810327	FR 1980-20590	19800925
FR 2465753	B1	19840427		
GB 2066824	A	19810715	GB 1980-30900	19800925
GB 2066824	B2	19830824		
US 4414280	A	19831108	US 1981-327062	19811203
US 4446269	A	19840501	US 1982-354194	19820303
PRIORITY APPLN. INFO.:			US 1979-79173	A2 19790926
			US 1980-176595	A 19800808

OTHER SOURCE(S): MARPAT 97:217423

AB Solvents for fluoropolymers useful in casting reverse osmosis membranes have the composition $\text{CF}_2\text{XCFXO}[\text{CF}_2\text{C}(\text{CF}_3)\text{FO}]_n(\text{CF}_2)_m\text{Y}$ (X = halogen; n = 0, 1; m = 1-3; Y = CO_2Me , SO_2F). Thus, 3276.1 g perfluoro[2-(2-fluorosulfonylethoxy)propyl vinyl ether] [16090-14-5] was chlorinated to give 2533.8g perfluoro[2-(2-fluorosulfonylethoxy)propyl-1,2-dichloroethyl ether] (I) [68860-43-5]. perfluoro[2-(2-fluorosulfonylethoxy-2-trifluoromethylethyl)]vinyl ether-tetrafluoroethylene copolymer [26654-97-7] (2 G) was dissolved in 45 g I, and 5 mL solution was cast to give a film which was dried at $80^\circ/300$ mm. The film was hydrolyzed with 28% NaOH at 80° to give a membrane which was tested in 0.3% NaCl in a hyperfiltration cell. The water flux d. at 5700 KPa was 1.872 ± 10^{-6} m/s, and the salt rejection was 82.6%.

IC C08K005-42; C08K005-10

INCL 524167000

CC 37-6 (Plastics Manufacture and Processing)

IT 4089-58-1

RL: RCT (Reactant); RACT (Reactant or reagent)
 (fluorination of)

IT 69116-73-0P 78010-39-6P

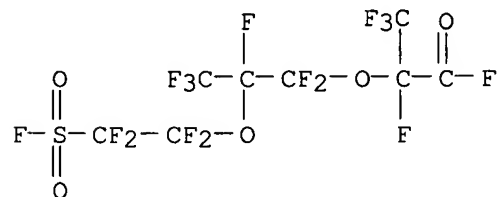
RL: PREP (Preparation)
 (preparation of)

IT 4089-58-1

RL: RCT (Reactant); RACT (Reactant or reagent)
 (fluorination of)

RN 4089-58-1 CAPLUS

CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,3,3,3-hexafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]propoxy]- (9CI) (CA INDEX NAME)



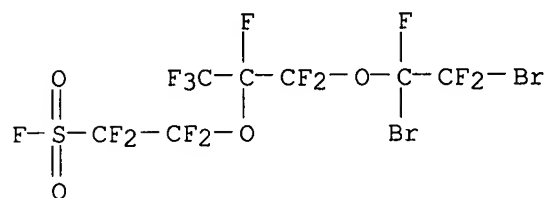
IT 78010-39-6P

RL: **PREP (Preparation)**

(preparation of)

RN 78010-39-6 CAPLUS

CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



L28 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1981:605062 CAPLUS

DOCUMENT NUMBER: 95:205062

TITLE: Solutions of copolymers of perfluoroethylene and a fluorosulfonated or carboxylated vinyl monomer in a saturated perhalogenated liquid

INVENTOR(S): Silva, Raimund Heinrich; Resnick, Paul Raphael; Smith, Roger Alton

PATENT ASSIGNEE(S): du Pont de Nemours, E. I., and Co., USA

SOURCE: Fr. Demande, 33 pp.

CODEN: FRXXBL

DOCUMENT TYPE: Patent

LANGUAGE: French

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FR 2465753	A1	19810327	FR 1980-20590	19800925
FR 2465753	B1	19840427		
US 4348310	A	19820907	US 1980-176595	19800808
PRIORITY APPLN. INFO.:			US 1979-79173	A 19790926
			US 1980-176595	A 19800808

AB C1CF2CC1FOCF2CF(CF3)OCF2CF2SO2F (I) [68860-43-5], C1CF2CC1FOCF2CF(CF3)OCF2CF2CO2Me [78010-35-2], FSO2CF2CF2OCF(CF3)CF2OCF(CF3)SO2F [78010-40-9], and 19 similar compds. are used as solvents for copolymers of F2C:CF2 and F2C:CFOCF2CF(CF3)OCF2CF2CO2Me or F2C:CFOCF2CF(CF3)OCF2CF2SO2F (II). The solns. are useful for the preparation and repair of membranes, for coating catalyst supports in the preparation of catalyst, etc. Thus, a solution of 2 g F2C:CF2-II copolymer [26654-97-7] in 45 g I was cast to prepare a membrane. The membrane was hydrolyzed with aqueous NaOH at 80° to prepare an ultrafiltration membrane which gave 82.6% rejection of NaCl during filtration.

IC C08F214-26; C08F002-06; B01D013-00; B01J035-00

CC 37-1 (Plastics Fabrication and Uses)

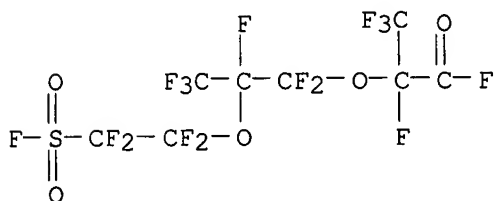
IT 4089-58-1

RL: **RCT (Reactant); RACT (Reactant or reagent)**
(decarbonylation of)

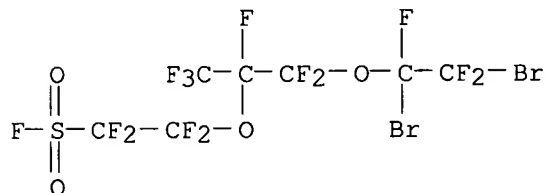
IT 27744-59-8P 78010-36-3P 78010-39-6P

RL: SPN (Synthetic preparation); **PREP (Preparation)**

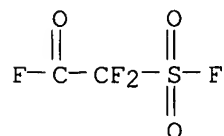
(preparation of)
 IT 677-67-8
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (reaction of, with tetrafluoroethylene)
 IT 4089-58-1
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (decarbonylation of)
 RN 4089-58-1 CAPLUS
 CN Propanoyl fluoride, 2,3,3,3-tetrafluoro-2-[1,1,2,3,3,3-hexafluoro-2-[1,1,2,2-tetrafluoro-2-(fluorosulfonyl)ethoxy]propoxy]- (9CI) (CA INDEX NAME)



IT 78010-39-6P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 78010-39-6 CAPLUS
 CN Ethanesulfonyl fluoride, 2-[1-[(1,2-dibromo-1,2,2-trifluoroethoxy)difluoromethyl]-1,2,2,2-tetrafluoroethoxy]-1,1,2,2-tetrafluoro- (9CI) (CA INDEX NAME)



IT 677-67-8
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (reaction of, with tetrafluoroethylene)
 RN 677-67-8 CAPLUS
 CN Acetyl fluoride, difluoro(fluorosulfonyl)- (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)



Search history

Keys 10/795995

02/24/2006

=> d his full

TRANSCRIPT FOR SERIAL NUMBER 10/795995 BEGINS WITH L18

FILE 'REGISTRY' ENTERED AT 15:36:34 ON 24 FEB 2006
L18 STRUCTURE UPLOADED

L19 1 SEA SSS SAM L18

D SCA

L20 11 SEA SSS FUL L18

SAVE TEMP L20 KEYSFLUSTRA/A

FILE 'CAPLUS' ENTERED AT 15:38:46 ON 24 FEB 2006
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FILE 'REGISTRY' ENTERED AT 15:39:07 ON 24 FEB 2006

D SCA L20

L*** DEL 6 S L20 (L) (RACT OR RCT OR RGT)/RL

FILE 'CAPLUS' ENTERED AT 15:43:53 ON 24 FEB 2006
L22 88 SEA ABB=ON PLU=ON L20 (L) (RACT OR RCT OR RGT)/RL

FILE 'REGISTRY' ENTERED AT 15:56:27 ON 24 FEB 2006

L23 STRUCTURE UPLOADED

L24 0 SEA SSS SAM L23

L25 6 SEA SSS FUL L23

SAVE TEMP KEYSFLUSTRB/A L25

FILE 'CAPLUS' ENTERED AT 15:57:59 ON 24 FEB 2006
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L27 18 SEA ABB=ON PLU=ON L25 (L) PREP/RL

L28 8 SEA ABB=ON PLU=ON L22 AND L27

FILE 'CASREACT' ENTERED AT 15:59:31 ON 24 FEB 2006
L29 1 SEA ABB=ON PLU=ON L20/RRT (L) L25/PRO
D SCA

FILE 'CASREACT' ENTERED AT 16:02:52 ON 24 FEB 2006
D STAT QUE L29
D IBIB ABS HIT L29 1

FILE 'REGISTRY' ENTERED AT 16:05:39 ON 24 FEB 2006

FILE 'CAPLUS' ENTERED AT 16:05:42 ON 24 FEB 2006
D STAT QUE L28
D IBIB ABS HITIND HITSTR L28 1-8

FILE HOME

FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file
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STRUCTURE FILE UPDATES: 22 FEB 2006 HIGHEST RN 874945-83-2

DICTIONARY FILE UPDATES: 22 FEB 2006 HIGHEST RN 874945-83-2

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TSCA INFORMATION NOW CURRENT THROUGH January 6, 2006

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*
* The CA roles and document type information have been removed from *
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* effective March 20, 2005. A new display format, IDERL, is now *
* available and contains the CA role and document type information. *
*

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FILE LAST UPDATED: 23 Feb 2006 (20060223/ED)

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<http://www.cas.org/infopolicy.html>

FILE STNGUIDE
FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Feb 17, 2006 (20060217/UP).

FILE BEILSTEIN
FILE LAST UPDATED ON JANUARY 17, 2006

FILE COVERS 1771 TO 2005.
FILE CONTAINS 9,428,406 SUBSTANCES

>>>PLEASE NOTE: Reaction Data and substance data are stored in separate documents and can not be searched together in one query. Reaction data for BEILSTEIN compounds may be displayed immediately with the display codes PRE (preparations) and REA (reactions). A substance answer set retrieved after the search for a chemical name, a compounds with available reaction information by combining with PRE/FA, REA/FA or more generally with RX/FA. The BEILSTEIN Registry Number (BRN) is the link

between a BEILSTEIN compound and belonging reactions. For more detailed reaction searches BRNs can be searched as reaction partner BRNs Reactant BRN (RX.RBRN) or Product BRN (RX.PBRN).<<<

>>> FOR SEARCHING PREPARATIONS SEE HELP PRE <<<

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* ARE BASED ON THE HIGHEST PRICE CATEGORY. THEREFORE; THESE *
* ESTIMATES MAY NOT REFLECT THE ACTUAL COSTS. *
* FOR PRICE INFORMATION SEE HELP COST *

NEW

* PATENT NUMBERS (PN) AND BABS ACCESSION NUMBERS (BABSAN) CAN NOW BE SEARCHED, SELECTED AND TRANSFERRED.
* NEW DISPLAY FORMATS ALLREF, ALLP AND BABSAN SHOW ALL REFERENCES, ALL PATENT REFERENCES, OR ALL BABS ACCESSION NUMBERS FOR A COMPOUND AT A GLANCE.

FILE BABS

FILE LAST UPDATED: 10 JAN 2006 <20060110/UP>
FILE COVERS 1980 TO DATE.

FILE CASREACT

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FILE CONTENT:1840 - 19 Feb 2006 VOL 144 ISS 8

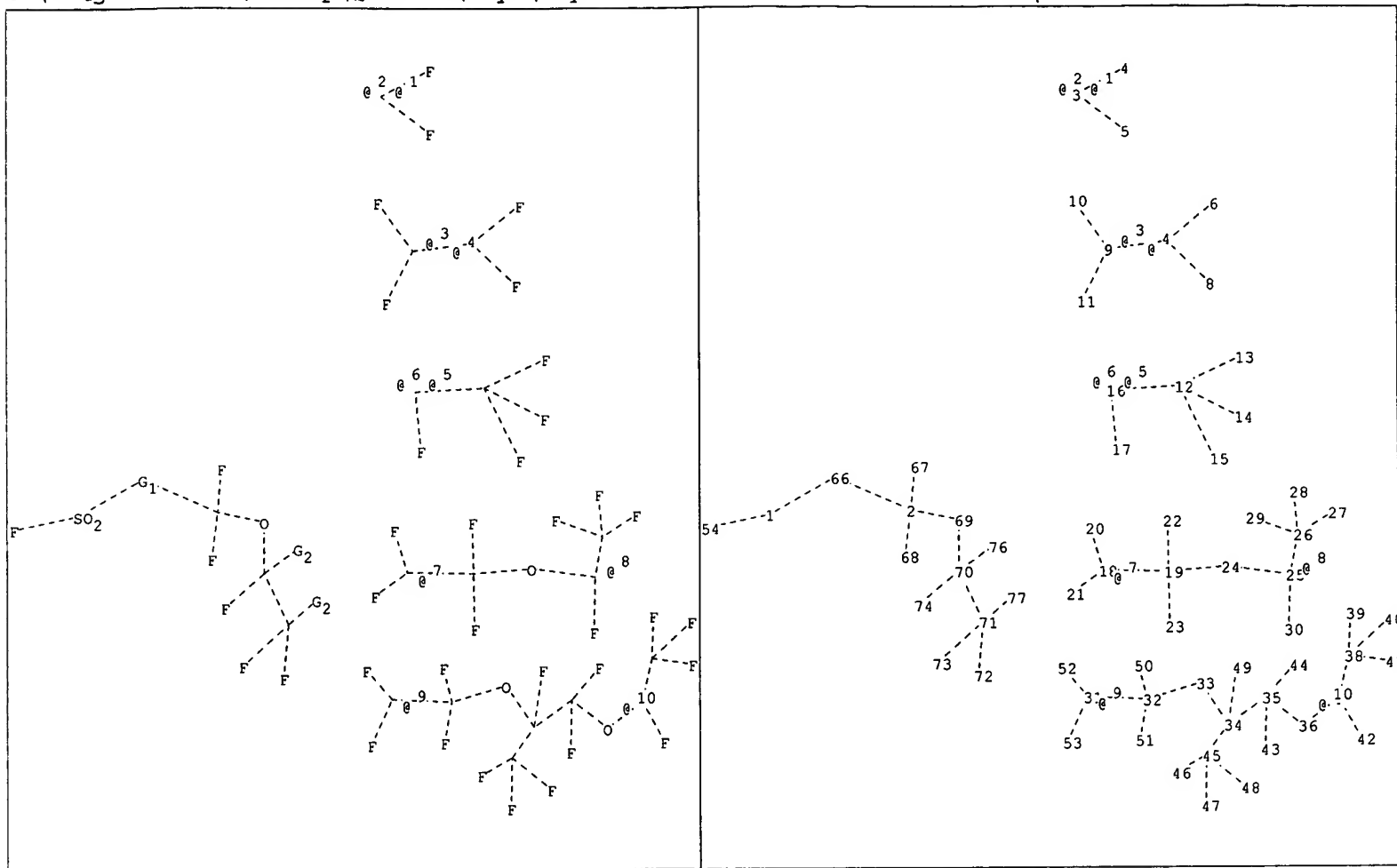
New CAS Information Use Policies, enter HELP USAGETERMS for details.

* *
* CASREACT now has more than 10 million reactions *
* *

Some CASREACT records are derived from the ZIC/VINITI database (1974-1991) provided by InfoChem, INPI data prior to 1986, and Biotransformations database compiled under the direction of Professor Dr. Klaus Kieslich.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=>



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25
 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47
 48 49 50 51 52 53 54 66 67 68 69 70 71 72 73 74 76 77

chain bonds :

1-54 1-66 2-66 2-67 2-68 2-69 3-4 3-5 6-7 7-8 7-9 9-10 9-11 12-13 12-14
 12-15 12-16 16-17 18-19 18-20 18-21 19-22 19-23 19-24 24-25 25-26 25-30 26-27
 26-28 26-29 31-32 31-52 31-53 32-33 32-50 32-51 33-34 34-35 34-45 34-49 35-36
 35-43 35-44 36-37 37-38 37-42 38-39 38-40 38-41 45-46 45-47 45-48 69-70 70-71
 70-74 70-76 71-72 71-73 71-77

exact/norm bonds :

1-54 1-66 2-66 2-67 2-68 2-69 3-4 3-5 6-7 7-8 7-9 9-10 9-11 12-13 12-14
 12-15 12-16 16-17 18-19 18-20 18-21 19-22 19-23 19-24 24-25 25-26 25-30 26-27
 26-28 26-29 31-32 31-52 31-53 32-33 32-50 32-51 33-34 34-35 34-45 34-49 35-36
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 70-74 70-76 71-72 71-73 71-77

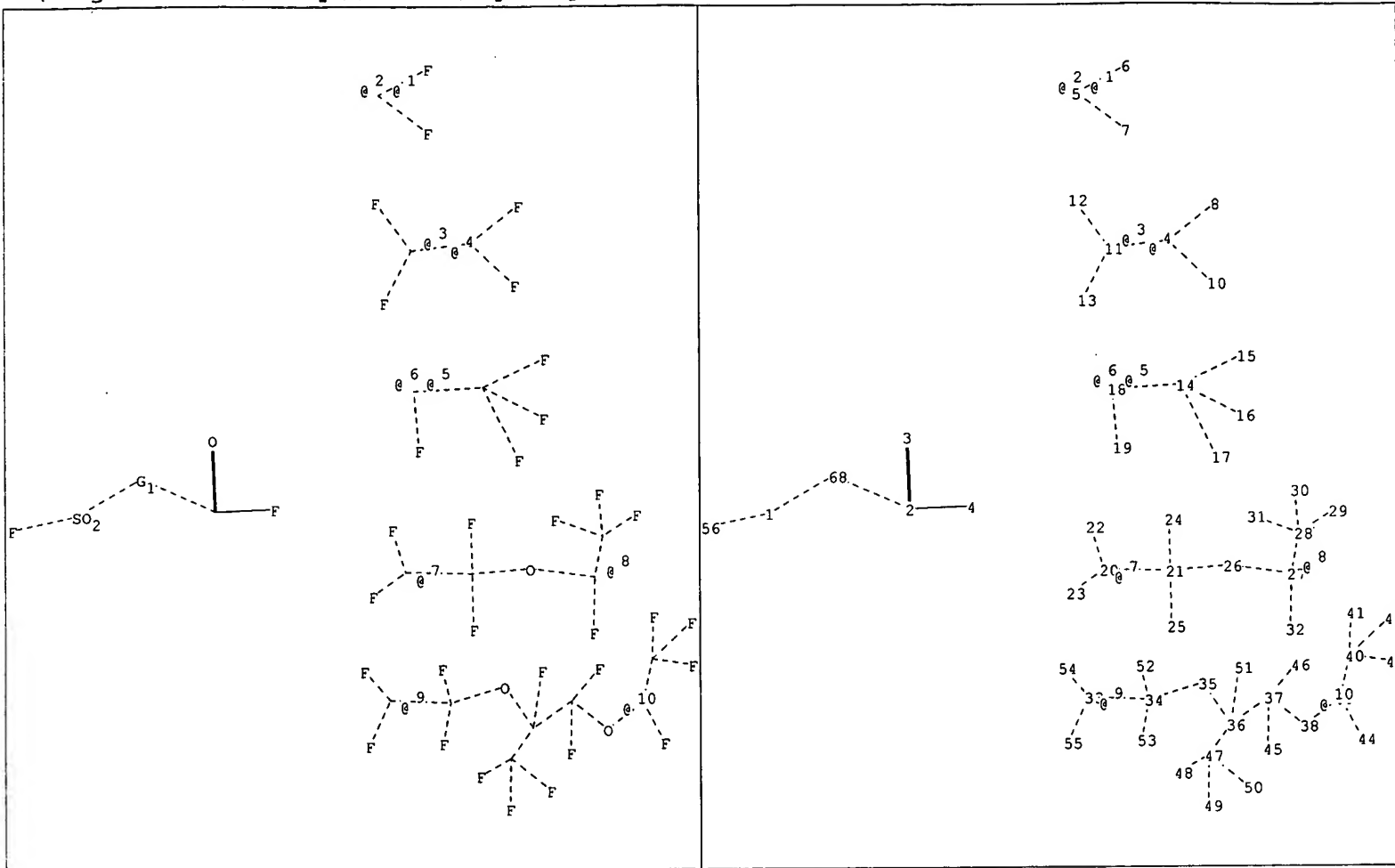
G1: [*1-*2], [*3-*4], [*5-*6], [*7-*8], [*9-*10]

G2: Cl, Br

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS
 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS
 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS
 29:CLASS 30:CLASS 31:CLASS 32:CLASS 33:CLASS 34:CLASS 35:CLASS 36:CLASS 37:CLASS
 38:CLASS

39:CLASS	40:CLASS	41:CLASS	42:CLASS	43:CLASS	44:CLASS	45:CLASS	46:CLASS
47:CLASS	48:CLASS	49:CLASS	50:CLASS	51:CLASS	52:CLASS	53:CLASS	54:CLASS
55:CLASS	56:CLASS	57:CLASS	58:CLASS	59:CLASS	60:CLASS	61:CLASS	62:CLASS
63:CLASS	64:CLASS	65:CLASS	66:CLASS	67:CLASS	68:CLASS	69:CLASS	70:CLASS
71:CLASS	72:CLASS	73:CLASS	74:CLASS	75:CLASS	76:CLASS	77:CLASS	



chain nodes :

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25
 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47
 48 49 50 51 52 53 54 55 56 68

chain bonds :

1-56 1-68 2-3 2-4 2-68 5-6 5-7 8-9 9-10 9-11 11-12 11-13 14-15 14-16 14-17
 14-18 18-19 20-21 20-22 20-23 21-24 21-25 21-26 26-27 27-28 27-32 28-29 28-30
 28-31 33-34 33-54 33-55 34-35 34-52 34-53 35-36 36-37 36-47 36-51 37-38 37-45
 37-46 38-39 39-40 39-44 40-41 40-42 40-43 47-48 47-49 47-50

exact/norm bonds :

1-56 1-68 2-3 2-68 5-6 5-7 8-9 9-10 9-11 11-12 11-13 14-15 14-16 14-17
 14-18 18-19 20-21 20-22 20-23 21-24 21-25 21-26 26-27 27-28 27-32 28-29 28-30
 28-31 33-34 33-54 33-55 34-35 34-52 34-53 35-36 36-37 36-47 36-51 37-38 37-45
 37-46 38-39 39-40 39-44 40-41 40-42 40-43 47-48 47-49 47-50

exact bonds :

2-4

G1: [*1-*2], [*3-*4], [*5-*6], [*7-*8], [*9-*10]

Match level :

1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 7:CLASS 8:CLASS 9:CLASS 10:CLASS
 11:CLASS 12:CLASS 13:CLASS 14:CLASS 15:CLASS 16:CLASS 17:CLASS 18:CLASS 19:CLASS
 20:CLASS 21:CLASS 22:CLASS 23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS
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 38:CLASS 39:CLASS 40:CLASS 41:CLASS 42:CLASS 43:CLASS 44:CLASS 45:CLASS 46:CLASS
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 56:CLASS

CASREACT

Keys 10/795995

02/24/2006

> file casreact

FILE 'CASREACT' ENTERED AT 16:02:52 ON 24 FEB 2006
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FILE CONTENT:1840 - 19 Feb 2006 VOL 144 ISS 8

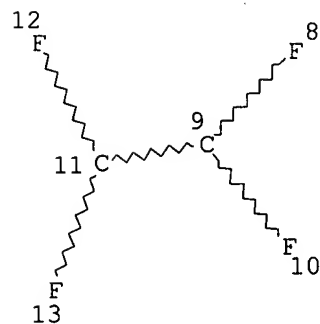
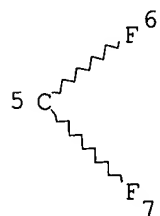
New CAS Information Use Policies, enter HELP USAGETERMS for details.

*
* CASREACT now has more than 10 million reactions *
*

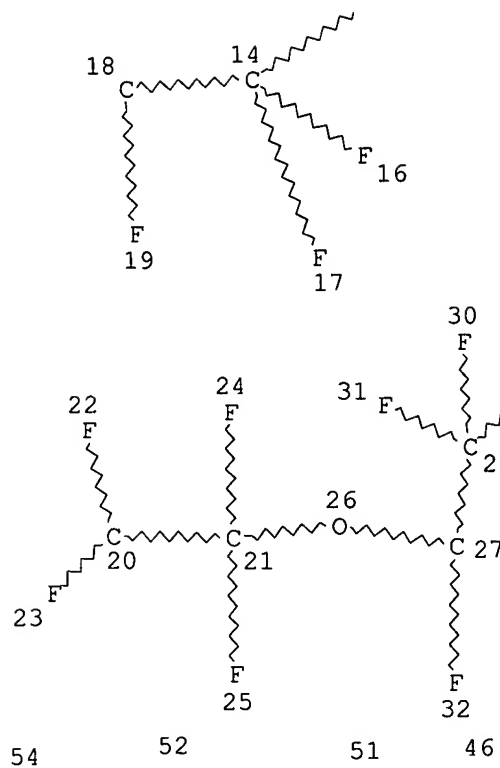
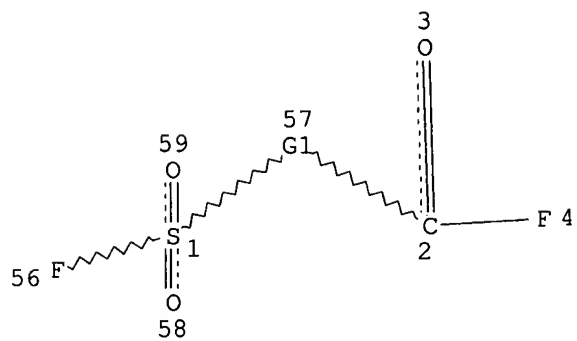
Some CASREACT records are derived from the ZIC/VINITI database (1974-1991) provided by InfoChem, INPI data prior to 1986, and Biotransformations database compiled under the direction of Professor Dr. Klaus Kieslich.

This file contains CAS Registry Numbers for easy and accurate substance identification.

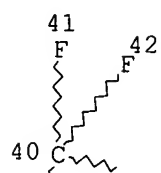
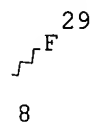
=> d stat que L29
L18 STR



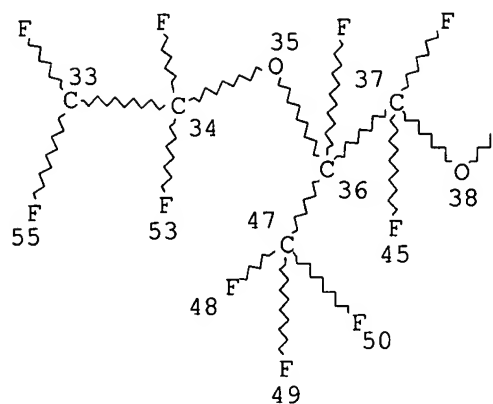
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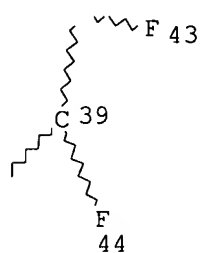
Page 2-A



Page 2-B



Page 3-A



Page 3-B

VAR G1=5-1 5-2/11-1 9-2/18-1 18-2/20-1 27-2/33-1 39-2

NODE ATTRIBUTES:

NSPEC	IS C	AT	1
NSPEC	IS C	AT	2
NSPEC	IS C	AT	3
NSPEC	IS C	AT	4
NSPEC	IS C	AT	5
NSPEC	IS C	AT	6
NSPEC	IS C	AT	7
NSPEC	IS C	AT	8
NSPEC	IS C	AT	9
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NSPEC	IS C	AT	27
NSPEC	IS C	AT	28

NSPEC IS C AT 29
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 NSPEC IS C AT 59

DEFAULT MLEVEL IS ATOM

MLEVEL IS CLASS AT 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17
 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38
 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 58 59

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

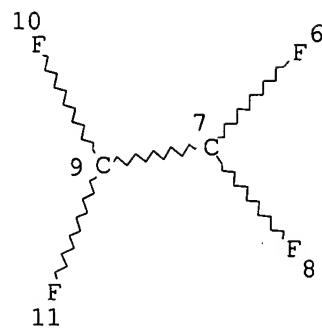
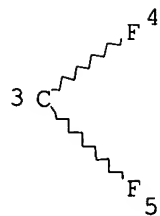
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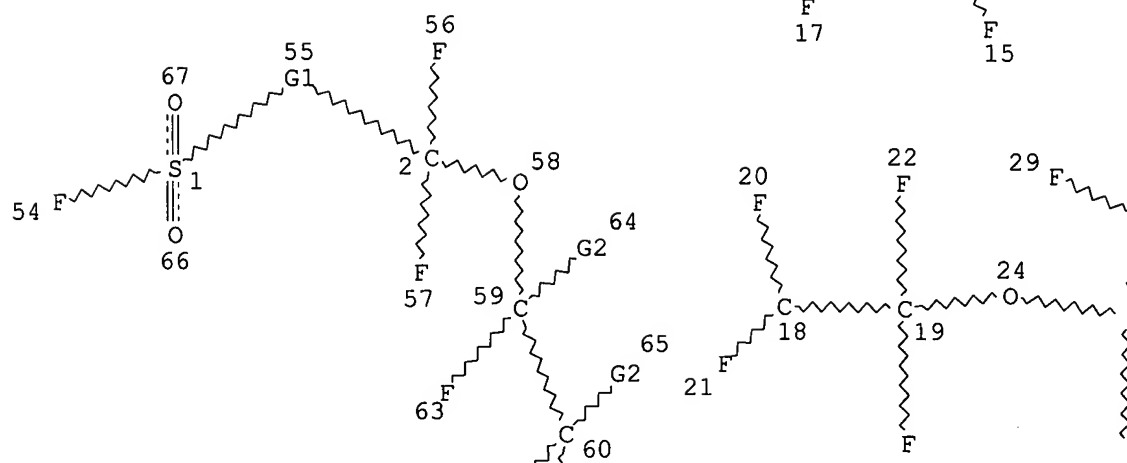
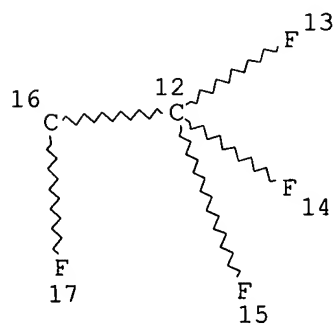
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L23 STR

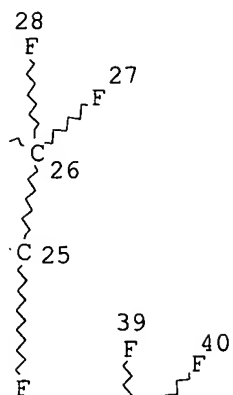
Cl 68Br 69



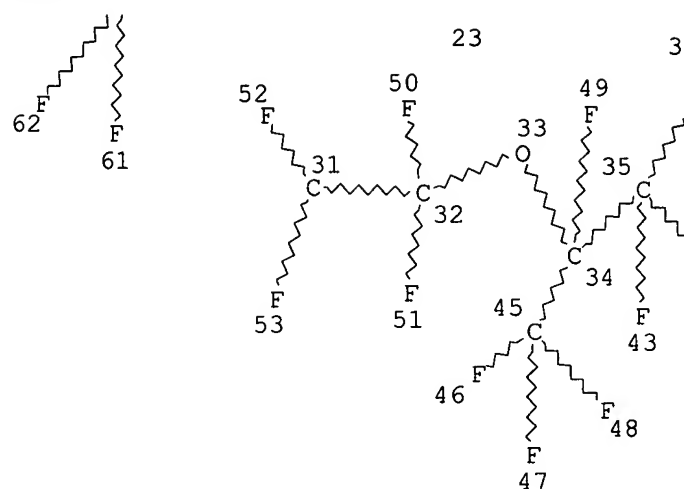
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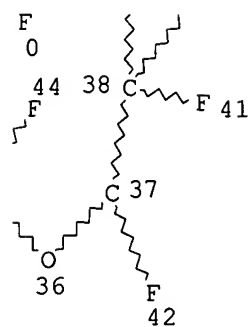
Page 2-A



Page 2-B



Page 3-A



Page 3-B

VAR G1=3-1 3-2/9-1 7-2/16-1 16-2/18-1 25-2/31-1 37-2

VAR G2=68/69

NODE ATTRIBUTES:

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NSPEC	IS C	AT	3
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NSPEC IS C AT 64
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MLEVEL IS CLASS AT 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17
18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38
39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 56 57 58 59 60
61 62 63 66 67 68 69
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 69

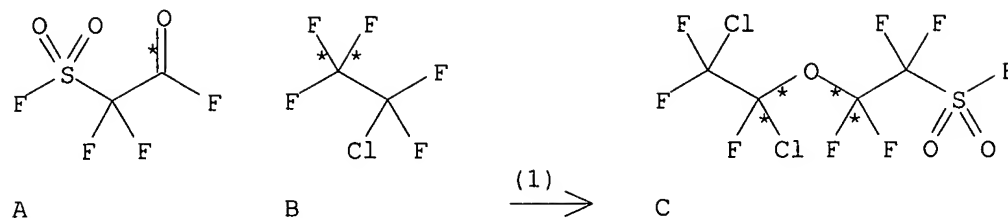
STEREO ATTRIBUTES: NONE
L25 6 SEA FILE=REGISTRY SSS FUL L23
L29 1 SEA FILE=CASREACT ABB=ON PLU=ON L20/RRT (L) L25/PRO

=> d ibib abs hit L29 1

L29 ANSWER 1 OF 1 CASREACT COPYRIGHT 2006 ACS on STN
ACCESSION NUMBER: 141:260266 CASREACT
TITLE: Process for preparing (per)fluorohalogen ethers by the
reaction of acyl fluorides with halogenated
1,2-difluoroethylenes
INVENTOR(S): Tortelli, Vito; Calini, Pierangelo; Millefanti,
Stefano
PATENT ASSIGNEE(S): Solvay Solexis S.p.A., Italy
SOURCE: Eur. Pat. Appl., 8 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1457484	A1	20040915	EP 2004-4344	20040226
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2004269535	A2	20040930	JP 2004-65994	20040309
US 2004199009	A1	20041007	US 2004-795995	20040310
CN 1539818	A	20041027	CN 2004-10033085	20040311
PRIORITY APPLN. INFO.:			IT 2003-MI444	20030311

OTHER SOURCE(S): MARPAT 141:260266
AB A process for preparing (per)fluorohalogen ethers containing the sulfonyl
fluoride group FSO2RCF2OCAFCAlF2 [A, Al = Cl, Br; R = (per)fluorinated
optionally containing one or more oxygen atoms] is described which comprises
the reaction of acyl fluorides FSO2RCOF in the liquid phase with elemental
fluorine and with olefinic compds. CAF:CA1F at -120° to
-20°, optionally in the presence of a solvent inert under the
reaction conditions.
REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

RX(1) OF 1 **A** + **B** ==> **C**

RX(1) RCT A **677-67-8**, B 76-15-3
 PRO C **144728-59-6**
 SOL 76-15-3 Ethane, chloropentafluoro-
 CON SUBSTAGE(2) 3 hours